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FRL ltr dtd 3 Sep 1971; FRL ltr dtd 3 Sep 1971



TECHNICAL REPORT 2179

CHARACTERISTICS OF EXPLOSIVE SUBSTANCES FOR APPLICATION IN AMMUNITION

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MAY 1955

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ORDNANCE PROJECT TA3-5002
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CHARACTERISTICS OF EXPLOSIVE SUBSTANCES FOR APPLICATION IN AMMUNITION

by

Alfred M. Anzalone James E. Abel Arthur C. Forsyth

May 1955

Picatinny Arsenal Dover, N. J.

Technical Report 2179

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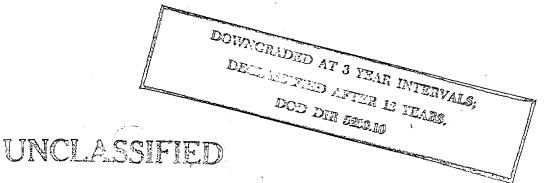
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OBJECT

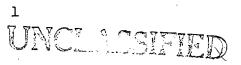
To record experimental data obtained on the properties of new and standard explosive compounds and compositions.

ABSTRACT

This project was established to provide for the testing and evaluation of new explosives synthesized in this Laboratory or on contract. During the past two years, the following materials were characterized: dinitrobenzfuroxan (DNBF), cesium DNBF, potassium DNBF, rubidium DNBF, mercuric DNBF, silver DNBF, lead DNBF; copper chlorotetrazole; stannous methylene bis (nitroso hydroxylamine) (i.e., stannous MEDNA), cupric MEDNA, lead MEDNA, mercury MEDNA, calcium MEDNA, cadmium MEDNA, barium MEDNA, and thallium MEDNA; lead nitratebis basic-4.6-dinitro-ortho-cresylate monohydrate; tris (ethylenediamine) chromic perchlorate; cyclotrimethylenetrinitrosamine (R salt): 2,4,6 trinitrotolyl 3 methylnitramine; cuprous thiocyanate, cuprous thiocyanate (29.1%) + potassium chlorate (70.9%); silver thiocyanate, silver thiocyanate (39.8%) +potassium chlorate (60.2%); lead thiocyanate, lead thiocyanate (53.5%) → potassium chlorate (46.5%); and silver cyanamide.

As a comparison, several specification grade military explosives were retested. These were: mercury fulminate; normal lead styphnate; lead azide; diazodinitrophenol; PETN; tetracene; Cyclonite; Haleite; TNT; Composition A-3; and tetryl. Some effort has also been devoted to the improvement of small-scale laboratory tests. The evaluation tests considered standard at this Arsenal for characterizing initiating type explosives left much to be desired. A need existed for a means of measuring properties such as the ignition time, burning time, linear length of the flame, ignition temperature, and shock wave impulse. This study is being continued to develop better techniques for measuring the properties of initiating materials, under Project TA3-5101, in an effort to obtain ideal explosives for optimum explosive trains.

To determine the above physical properties, two new instruments were developed, a squib tester, and an ignition temperature apparatus.





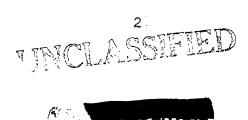


INTRODUCTION

1. This report presents the data obtained in the characterization of explosives which have been synthesized under projects controlled by this laboratory or have appeared promising for application in ammunition. Many tests accepted as standard at Picatinny Arsenal were used in this work. However, these tests were not sufficient to determine the possible military use of new primary explosives. Therefore, part of the work of this project was devoted to the development of a squib tester and an ignition temperature apparatus to permit the measuring of such properties as: ignition temperature, ignition time, duration of flame, linear length of flame, and shock wave impulse. In addition, the characteristics of the particles of explosive were obtained by photmicrographic techniques.

DISCUSSION

- 2. The compounds which were evaluated more fully are mercury fulminate, dextrinated lead azide, normal lead styphnate, diazodinitrophenol, Cyclonite, Haleite, PETN, Composition A-3, tetryl, and tetracene. The recently prepared compounds which were evaluated are: silver cyanamide, several metal salts of methylene-bis (nitroso hydroxylamine), dinitrobenzfuroxan (DNBF) and various salts, copper chlorotetrazole, and a few of the metal salts of thiocyanic acid, by itself, and in combination with potassium chlorate. (Tables 1-5).
- 3. Included in the evaluation are tests for impact sensitivity, explosion temperature, heat stability, hygroscopcity, water solubility, heats of explosion and combustion, gas volume, crystal density, crystal size and color, brisance, friction sensitivity, specific heat, power, plate dent test, and rate of detonation.
- 4. Coincident with the evaluation new compounds, work was started on the development of new small-scale tests to make possible a more complete evaluation of the explosives. Since a number of the new compounds were expected to be used as primary explosives, it was decided to study the characteristics of the flame emitted by the burning explosive. The Instrumenta tion Unit of Picatinny Arsenal designed and constructed a firing device (Fig 1) which enables the operator to control



the current input through the wire bridge of a measured resistance, by means of variable controls, from 0 to 2 amperes and to register the resistance of the squib from 0 to 11 ohms. It is also possible to apply a specific current for a definite time interval. Outlets were provided for connecting a chronograph and a photocell which are used in recording ignition time.

- 5. When the current (I), resistance (R), and ignition time (t) are known, the energy input (E) required to ignite any explosive may be calculated by the formula: $E = I^2Rt$ (107) where E is expressed in ergs, I in amperes, T in seconds, and R in ohms. This formula was used to determine the energy values reported in Table 6 for ignition of the various explosives in bridge-wire squibs.
- 6. Since carbon-bridge-type squibs are also used in several fuze designs, tests similar to those made with the wire-bridge-type squibs were run. The firing energies needed for this type of initiation were calculated from the results obtained, and are listed in Table 6 (See also Fig 2).
- 7. The burning times, or the duration of flame produced in the visible bands of the spectrum, were measured with the equipment shown in Figure 1. The results (Table 6) show that the lead salt of DNBF has the longest visible duration of flame, while the alkali salts do not register on the photocell at all.
- 8. Although this test gave information on the characteristics of the flame, it was believed that photographs would permit a more comprehensive study of the flame structure. Such pictures were made using a technique similar to that used in obtaining the burning times. In this test the fixture was placed in a dark room in front of a plate camera. The camera shutter was opened before firing and remained open until the squib had been fired.
- 9. Figures 3,4, and 5 show the flames emitted by several of the compounds and mixtures tested. The normal lead styphnate produces a much larger flame than colloidal normal lead styphnate. Mercury fulminate also produces a large flame, while dextrinated lead azide gives a small spit of flame. (Grouping some of these compounds according to their

use, the FA70 mix is a percussion primer mixture, while the modified 7L and the PA 100 are stab type mixtures.) The results obtained from DNBF salts and silver cyanamide corroborate those obtained from the duration of the flame test. Again, the lead DNBF registered a flame which the alkali salts did not. An interesting observation in the case of the lead DNBF is that the flame emitted was much larger than that of any of the other mono-component explosives.

- 10. From the results tabulated in Table 6, it can be observed that the addition of an oxidant to an explosive or a fuel in stoichiometric quantities increases the burning rate and decreases burning time. By controlling the amount of added oxygen, from zero percent to stoichiometric amounts, it was believed that the length of the flame could also be controlled. An attempt to illustrate this is shown in Figure 5. Here, the cesium styphnate alone (B), having a negative oxygen balance, has a flame three times as long as that produced when it is combined with potassium chlorate (C). Since the presence of sufficient oxygen permits a much more rapid rate of burning, the duration of the visible flame is much shorter. It can be observed (Fig 5), that the length of the flame is similarly shorter.
- ll. In the case of silver cyanamide (D), a fuel, ignition did occur but no visible flame was observed or recorded (See Table 6 and Figure 5-D). By adding potassium chlorate, the flame (Table 6) was made visible to the photocell and was recorded by the camera (Figure 5-E).
- 12. The shock wave emitted in an open air system, was measured for several compounds (Fig 6). Figure 7 shows the equipment which was used to record the character of the shock wave produced by the explosive when fired in an MlAl squib and received by a piezoelectric crystal gage at a distance of six inches from the squib being tested. The oscillographs of the resulting shock waves are shown in Figure 6. A comparison of the results shows the potassium DNBF to have a lower peak pressure than the other two alkali salts. These two salts, rubidium and cesium, have peak pressures similar to basic lead styphnate, which is used to initiate lead azide in some low-energy electric-type detonators. It is of interest to note that the lead DNBF which produced such a large flame

did not emit a shock wave sufficiently strong to be recorded.

13. The Fisher sub-sieve sizer was considered too hazardous to be used in determining the particle size of these explosives. Therefore, an alternate method was used in which photomicrographs were taken and the average particle size or crystal size measured by a superimposed scale.

EXPERIMENTAL PROCEDURE

- 14. The characterization tests used in this work, except for those referred to in detail in this report, are described in Picatinny Arsenal Technical Report 1401 (Revision 1).
- 15. Those explosives which are described as military explosives have been prepared commercially according to specifications and were given a conformatory analysis before they were used. All new materials investigated for use as explosives were prepared as laboratory samples were of the highest purity possible.
- 16. The following procedure was used in determining ignition time, burning time, and flame characteristics. The MIAL squib was filled with a weighed amount of the material to be tested, placed in a mount, connected to the firing leads from the squib tester, and set in front of the photocell in a hood protected from any outside light (See Fig 1). Next the resistance of the bridge was determined with a potentiometer, the desired input current set on the dial, and the squib fired by pressing the firing button on the face of the squib tester. This actuated the chronograph, which was stopped by the resulting flash of the material firing in front of the photocell. The ignition time was read in milliseconds, the chronograph having an accuracy of 0.1 millisecond.
- 17. A General Electric time interval meter (G. E. catalog No. 5106917) and a phototube preamphifier (G. E. catalog No. 51115576) were used to record the burning times (flame duration) of the compounds tested. The compound to be tested was loaded into an MIA1 squib which was placed 1-1/2 inches away from the phototube in a standard hood protected from the light by a dark curtain (Photo 1). When the resistance was determined, the desired input current set on the dial, and the time interval meter set on open, the

squib was fired. The burning time in milliseconds was then read from the dial. The measured time is the total length of the visible burning time from zero start back to zero finish.

- 18. The flame characteristics of the compounds were determined by taking photographs of the flames in a darkened walk-in hood. The MIAl squib was first filled with a definite weight of the material and placed in a clamp in front of a six-inch white ruler held vertically against a black background. A 35 mm camera equipped with a Leica lens was used with the lens fully opened (f/3.5). Kodak Super XX film was used in the camera, which was placed three feet from the squib. The squib was fired using the squib tester set at 600 milliamperes of current.
- 19. The results obtained for the shock wave tests were picked up with a piezoelectric crystal gage using a barium titanate crystal and recorded by an oscilloscope camera apparatus. The MIAl squib was first loaded by hand tamping, with a definite weight of material and placed two inches away from and parallel to the face of the piezo crystal gage. This gage was then connected to an oscilloscope (Dumont Model 304H-Serial No. 2399) and the resulting oscillogram photographed on linagraph film.
- 20. The energy to initiate explosives in carbon-bridge-type detonators was determined using the T18E4 detonator squib assembly. These squibs were fired by passing 110 volts AC through an AC to DC converter, to a variable condenser, and then through the firing leads to the T18E4 holder. An ohmmeter was used to check the resistance of the detonator before firing, and to set the firing voltage. The maximum voltage used was three hundred volts. The variable condenser (Model No. 83595-Central Scientific Co.) has two dials for setting the charge; one ranges from 0.01 to 0.10 mfd and the other from 0.1 to 1.0 mfd. Hence, by keeping the resistance constant and varying the voltage and capacitance, it is possible to control the energy used to initiate the explosive. This energy can be calculated by using the formula \underline{E} 5cv², where \underline{E} is in ergs, C is in microfarads, and v is in volts.

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- B: Tomlinson, W. R., Jr, <u>Properties of Explosives of Military Interest</u>, Picatinny Arsenal Technical Report 1740, June 1949.
- C. Gaughran, R. J., Abel, J. E., and Forsyth, A. C., <u>Development of Optimum Explosive Trains</u>, <u>Picatinny</u> Arsenal Report MR 43.
- D. Abel, Alster, Forsyth, Jackson, and Pataky, <u>Development of A New Ignition Temperature Apparatus</u>, Picatinny Arsenal Technical Report 2093, November 1954.

INCLOSURES

Tables 1-6
Figures 1-13

TABLE 1

Military Explosives Used as or in Primers Compared With New Explosives Having Similar Properties

	Mercury Fulminate	Normal Lead Styphnate	Dinitro- Benz- Furoxan	Cesium DNBF	Potassium DNBF	Rubidium DNBF	Mercuric DNBF	Silver DNBF	Cupric DNBF	Lead DNBF (hydrate)
Impact Sensitivity, P. A. Machine										
2 kg. wt, inches 1 lb. wt, inches average chge, wt, grams Explosion Temperature, OC	2° 3° 0.030° 210°	36 33 0.021 282 ⁰	9 # 0.010 310	2 6 0.012 210	3 6 0.007 250	3 7 0.009 225	3 14 0.013 215ª	4 13 0.10 2008	4 40 0.10 260	7 32 0.020 185
100°C Heat Test, % Loss 1st 48 hrs 2nd 48 hrs. Explosion, 100 hrs.	exploded ^c 16 hrs	0.38 0.73 none	0.28 0.08 none	0.08 0.13 none	0.30 0.05 none	0.37 0.00 none	5.20 2.85 none	0.73 0.20 none	9.10 1.18 none	7.05 0.77 none
Hygroscopicity at 30°C, % Gain										
75% R.H. 90% R.H. Water Solubility at 30°C, gms/100 gm Heat of Explosion, cal/gm Heat of Combustion, cal/gm Cas Volume, cc Particle Size, Microns Color	0.00 0.02 0.042 427 938 243 554 gray ^c	0.00 0.02 0.074 457 1251 368 48 yellow to	0.14 0.16 * 1790 * golden- brown	1.70 3.45 0.312 * * yellow- orange	0.11 0.27 0.245 725 2209 604 74 orange to	0.62 1.77 0.239 * 1931 * violet	* 0.041 * # # red	0.28 0.31 0.284 698 1864 484 * brick-	8.92 4.96 0.017 * 1948 * black	* * 0.162 * * * red
Crystal form Absolute Density	rhombic ^c 4.43 ^c	hexagonal ^c 3.08 ^c	# 1.703	*	platinates 2.026	* 2.208	*	# 2.529	# #	*
200 gm Bomb Sand Test - gms. sand crushed when initiated by: Black Powder Fuze 300 mg, Lead Azide 200 mg. Lead Azide + 100 mg Tetryl	23.4° max w/BPF° max w/BPF°	11.4° 24.0° max w/L.A.°	* *	12.4 35.8 34.8	9.5 44.8 43.6	7.0 38.9 40.0	no fire 0.0 5.0	no fire 44.1 57.5	no fire 27.8 29.7	no fire 25.7 27.6
Friction Sensitivity Fiber Shoe Steel Shoe	detonates ^c	detonates	*	**	**	**	**	## ##	## ##	*

a. Smoked - no detonation

b. Flashed

c. Work reported previously

^{*} Further work suspended to enable the investigation of the more promising compounds to be completed first.

^{**} Since the friction sensitivity test used at this Arsenal has, as one of its limitations, the fact that the test also reacts to impact sensitivity; those compounds which show a very low impact value are assumed to give a positive test on the friction machine.

CONFIDENTAL TABLE 1 (Con't)

Military Explosives Used as or in Primers Compared With New Explosives Having Similar Properties

		Copper Chloro- Tetrazole	Stannous Methylene Bis (Nitroso Hydroxyla- mine)	Cupric MEDNA	Lead MEDNA	Mercury MEDNA	Calcium IMEDNA	Cadmium MEDNA	Barium MEDNA	Thallium MEDNA	Lead Nitrate Bis Basic- 4,6 Dinitro- Ortho-Cres- ylate Mono- hydrate
Impa	ct Sensitivity, P. A. Machine										
Expl	2 kg. wt, inches 1 lb. wt, inches average chgc, wt, grams osion Temperature, ^{QC}	1 3 0,009 305	1 3 0.015 385 ^b	4 42 0.011 190 ⁸	2 18 0.015 240 ^b	2 * 0.021 165°	10 * 0.090 295a	12 0,014 240ª	15 * 0.016 275	2 10 0.22 225	11 * 0.022 265
100°	C Heat Test, % Loss '										
	lst 48 hrs. 2nd 48 hrs. Explosion, 100 hrs.	2.67 0.10 none	0.57 2.17 none	4.35 1.23 none	3.45 0.08 none	34.15 2.37 none	1.92 0.00 none	1.25 0.17 none	10.68 0.00 none	0.18 0.00 none	0.50 0.00 none
Hygr	oscopicity at 30°C, % Gain										
	75% R.H.	*	0.00	2.24	0.00	0.00	0.00	4.87	7.64	0.16	0.26
	90% R. H.	3.11	0.00	4.87	0.00	0.00	39.23	5.36	9.58	0.18	0.77
Wate	r Solubility at 30°C, gms/100 gm	0.02	0.02	0.02	0.02	*	*	#	#	0.67	0.23
Heat	of Explosion, cal/gm	*	1012	975	612	*	*	*	*	*	*
	of Combustion, cal/gm	*	1134 '	1024	620	*	*	*	*	#	*
	Volume, cc	*	243	247	126	#	*	*	*	#	*
	icle Size, Microns	#	#	*	*	*	#	#	*	#	*
Colo		blue	cream	blue	buff white	buff- white	white	white	white	white	brown
	tal form	#	#	*	#	#	#	#	*	*	*
Abso	lute Density	2.040	2.772	#	*	*	*	* .	•	*	*
200	gm Bomb Sand Test — gms. sand crushed when initiated by:										
	Black Powder Fuze	17.0	2.1	failed		failed	#	failed	failed	*	1.2
	300 mg. Lead Azide	25.3	27.6	*	20.1	#	#	#	*	*	17.5
	200 mg. Lead Azide 100 mg Tetryl	27.4	26.5	*	17.8	#	*	#	#	*	17.1
Frict	tion Sensitivity										
	Fiber Shoe	\$ -\$	##	**	## ·	**	#	*	**	##	*
	Steel Shoe	**	**	**	**	##	*	*		**	#

a. Smoked - no detonation

b. Flashed

Work reported previously Further work suspended to enable the investigation

of the more promising compounds to be completed first.
** Since the friction sensitivity test used at this Arsenal has, as one of its limitations, the fact that the test also reacts to impact sensitivity; those compounds which show a very low impact value are assumed to give a positive test on the friction machine.

TABLE 2

Military Explosives Used As Intermediate Charges And New Explosives Having Similar Properties

	Lead Azide (dextri- nated)	Diazo- Dinitro- phenol	PETN	Tetracene	Hexamine Chromic Perchlo- rate	Tris- Ethylene Chromic Perchlo- rate
Impact Sensitivity, P. A. Machine						
2 kg. wt, inches 1 lb. wt, inches average chgc, wt, grams Explosion Temperature, OC	5* 23 0.027 340*	4 7 0.015 180	6* a 0.016* 225*	2* a 0.016* 160*	8 30 0.020 350	3 18 0.012 315
100℃ Heat Test, \$ loss						
lst 48 hrs. 2nd 48 hrs. Explosion, 100 hrs.	0.34* 0.05* none*	2.10* 2.20* none*	0.10* 0.00* none*	23.2* 3.4* none*	0.08 0.00 none	0.35 0.00 none
Hygroscopicity at 30℃, % Gain						
75% R.H. 90% R.H. Water Solubility at 30°C, gms/100 gm Heat of Explosion, cal/gm Heat of Combustion, cal/gm Gas Volume, cc Particle Size, Microns Color	0.00 0.48 0.005 342 630 340 554 gray*	0.00* 0.04* 0.040 820* 3243 865 28 greenish- brown *	0.00* 0.00* a 1370* 1960* a 192 white*	0.00 0.80* a 664* 2758 451 546 pale- yellow*	0.02 0.05 b 1040 1110 125 50 yellow	4.50 26.0 18.4 ** ** ** yellow
Crystal form	rhombic*	elongated*	needles*	needles*	**	**
Absolute Density 200 gm Bomb Sand Test - gms. sand crushed when initiated by:	4.68	1.63*	1.77*	a.	1.95	##
Black Powder Fuze 300 mg. Lead Azide 200 mg. Lead Azide + 100 mg Tetryl Priction Sensitivity	23,4 max w/BPF	45.6* 47.5* max w/L.A.*	0.00 62.7* max w/L.A.*	4.0* 28.2* max w/L.A.*	32.3 41.8 40.6	** ** **
Fiber Shoe Steel Shoe	detonates*	detonates	unaffected* cracka*	detonates	**	## ##

- a. The future use of this compound at Present seems doubtful and further characterization was not warranted.
- b. The sample reacted with the wafer to form a gelatinous precipitated. $% \left(1\right) =\left(1\right) \left(1\right) \left($
- * Results obtained by other investigators
- ** Results not obtained as yet

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TABLE 3 Standard Military Explosives Used As Base Charges, Boosters or in Main Charges, And New Explosives Having Similar Properties

	Cyclonite (RDX)	Haleite (EDNA)	TNT	Compo- sition A-3	Tetryl	Cyclotri- methylene Trinitros- amine	2,4,6, Trini- tro-Tolyl- 3-Methyl- nitramine
Impact Sensitivity, P.A. Machine							
2 kg. wt, inches average wt. of charge, grams Explosion Temperature, °C, 5 secs.	8* 0.018* 260*	14* 0.017* 189*	14* 0.017* 475*	16* 0.017* 250*	8* 0.017* 257*	8 0.010 220	10 0.013 235
100°C Heat Test, % Loss							
lst 48 hrs. 2nd 48 hrs. Explosion, 100 hrs.	0.04* 0.00* none*	0.2* 0.3* none*	0.2* 0.2* none*	0.2* 0.2* none*	0.07 0.07 none	8.79 2.98 none	0.55 0.37 none
100 °C Vacuum Stability Test							
Weight, grams Gas, mls Time, hrs Water Solubility at 30°C, gms/100 gms Hygrosopicity at 30°C, % Gain at 90% R.H. Heat of Explosion, cal/gm Heat of Combustion, cal/gm Gas Volume, cc Particle size, microns Color	5.0* 0.7* 40* 0.005* 0.00 1280* 2285* ** 388 white*	5.0* 0.5* 40* 0.005 0.01* 981* 2477* ** 192 white*	5.0* 0.10* 40* 0.0219* 0.00* 1060 3620 ** buff*	5.0 0.42 40 0.90* 0.00* ** 1210* ** 862 white to	5.0* 0.3* 40* 0.09 0.04* 1130* 2925* ** 605 yellow*	2.5 9.19 40 0.28 0.02 876 3158 1030 6.8 yellow	2.5 1.07 40 0.01 0.07 752 3323 938 12.5 buff
Crystal Form Crystal Density or Specific Gravity	ellipsical* l.82*	** 1.71*	flakes* 1.65*	buff* ellipsical* **	ellipsical* 1.73*	** 2.14	** 1.64
200 gm Bomb Sand Test - gms of sand crushed when initiated by:							·
Black Powder Fuze 300 mg Lead Azide 200 mg Lead Azide + 100 mg Tetryl	0.00* 60.2* max w/L.A.*	0.00* 52.3* max w/L.A.*	0.00* 48.3* max w/L.A.	0.00* 51.5* max w/L.A.*	0.00* 54.2* max w/L.A.*	0.00 59.2 54.1	0.00 55.0 49.0
Priction Sensitivity							
Fiber Shoe Steel Shoe	unaffected* exploded*	unaffected* unaffected*	unaffected* unaffected*	unaffected* unaffected*	unaffected* crackles*	unaffected unaffected	unaffected crackles
Bellistic Mortar (TNT = 100) Power No. of Trials	150* 100 *	139* 34*	100* standard	135* 11*	130* 51*	130 8	113 11
Plate Dent Test Method Condition Confinement Density, gms/cc Brisance, % (TNT = 100)	A* pressed* yes* 1.50* 1.35*	A* pressed* yes* 1.50* 122*	A* pressed* yes* 1.50* 100*	B* pressed* no* 1.61* 126*	A* pressed* yes* 1.50* 116*	44 44 44 46	** ** **
Rate of Detonation							
meters/second (av. of 10 trys) Density, gms/sec.	8795* 1.82*	8820* 1.71*	7187 * 1.65*	8200* 1.61*	7972* 1.73*	7055ª 1.48	6580 1.49

a. Difficulty in going high order; this result is not considered final
 bata previously reported by other investigators
 Information not available yet

TABLE 4

Puels, And Fuel-Oxidant Mixtures to Replace T61 Primer Mix

Compound	Explosion Temperature, OC(5 secs) Sample 10 mg		eat Test °C, % Lo 2nd 48 Hrs			uum Sta est, 10 Gas, mls	bility O ^O C Hrs No. Expl		mpact Ser T.P4.Al Wt. of Charge, gms	lachine 4 oz		30	city	Anal % Me Found	
Cuprous Thiocyanate	None	0•171	0.03	None	1.0	0.17	40	None	0,012	None	0.013		-	51.8	52.3
Cuprous Thiocyanate (29.1%) + Potassium Chlorate (70.9%)	240°	0.13	0.02	None	1.0	0.41	40	ı	0,013	15	0.012	0.06 const	1.83	-	-
Silver Thiocyanate	None	0.26	0.01	None	1.0	0.97	40	None	0.014	None	0.016	after -	400 hrs	64.5	65.0
Silver Thiocyanate (39.8%) + Potassium Chlorate (60.2%)	200°	0.11	0.01	None	1.0	0.43	40	1	0.015	14	0.017	0.01	0.00	-	-
Lead Thiocyanate	None	0.18	0.01	None	1.0	0.44	40	None	0.015	None	0.015	-	-	63.9	64.1
Lead Thiocyanate (53.5%) + Potassium Chlorate (46.5%)	205 °	0.06	0.01	None	1.0	0.27	40	1	0.016	13	0.014	0.02	0.001	-	-
Silver Cyanamide	245	0.18	0.00	None	-		-	40d	0.019	62 8	0.018	0.35	0.35	84.3	84.3

 $\underline{\mathbf{a}}$ One pound weight

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<u>TABLE 5</u>

Ignition Temperature of Explosive Compounds*

•	Point of Immediate Ignition, OC		st Point gnition Time in Seconds
Potassium Dintirobenzfuroxan	267	240	9.8
Basic Lead Styphnate	312	291	17.5
Copper Chlorotetrazole	348	312	4.2

* Ref. PATR 2093, Nov 1954

TABLE 6

Behavior of Standard and Experimental Primary Explosives in Items Electrically Initiated

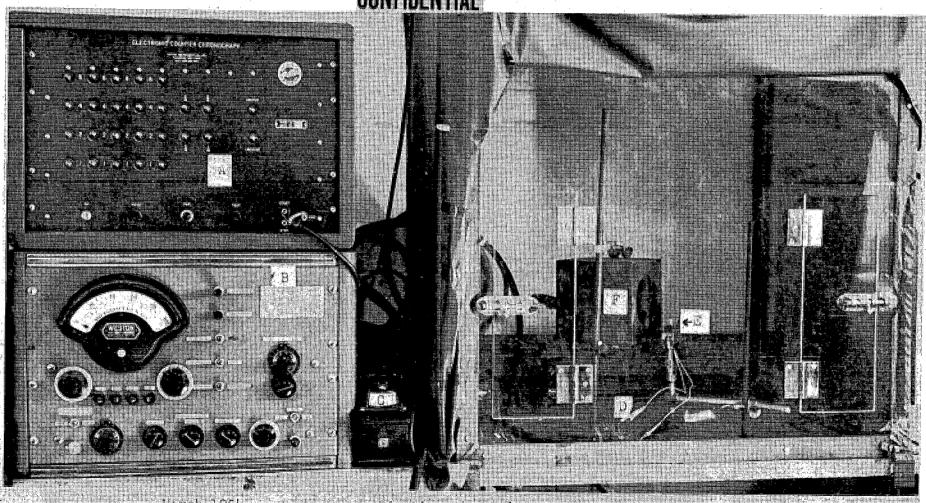
			Merca Fulmi				Norma Lead Styphr	Ī	
	Test No.	1	2	3	4	1	2	3	4
M1A1 Squib	Weight - Mgms. Resistance - Chms Current - Mamps Delay Time - Msecs Flame Duration - Msecs Energy - Ergs	50 1.07 350 75.95 19 96,300	50 1.05 400 35.36 20 59,500	50 1.05 500 19.16 22 50,400	50 1.07 600 13.50 21 52,050	65 1.14 400 34.02 4.4 62,000	65 1.12 500 15.05 5.5 42,300	65 0.98 600 11.09 6.0 39,200	65 1.08 700 8.90 4.8 47,100
	Test No.	1	2			1	2		
T18E4 Carbon Bridge	Volts Condenser - Mfd. Resistance - Ohms Energy - Ergs Remarks	300 .01 2600 0.0 No Fire	300 1.1 3400 0.0 No Fire			40 .01 2500 0.0 No Fire	50 .01 1500 125 Fired		
			Hexami Chromi Perchlo	ic		г	Rubidi initrobenz	_	
	Test No.	1	2	3	4	1	2	3	4
MlAl Squib	Weight - Mgms. Resistance - Ohms Current - Mamps. Delay Time - Msecs. Flame Duration - Msecs Energy - Ergs.	20.0 1.08 500 37.29 100+ 100,900	20.0 1.07 550 33.11 30 85,500	20.0 1.02 700 18.22 34 91,000	20.0 1.07 900 15.81 42 137,000	10 1.06 300 68.27 No Flash 65,200	10 1.01 550 6.90 No Flash 21,000	10: 1:03 600 5:75 No Flash 21,300	10 1.08 900 2.68 No Flash 23,400
	Test No.	1	2	3		1	2	3	
T18E4 Carbon Bridge	Volts Condenser - Mfd. Resistance - Ohms Energy - Ergs. Remarks	300 .01 6000 0.0 No Fire	300 1.1 5800 0.0 No Fire	300 1.1 5800 495,000 Fired with 10,000 psi		50 .01 1200 0.0 No Fire	100 .01 2400 5.0 Fired at 12,000 psi	300 .01 2900 450 Violent	

TABLE 6 (Con't)

		I	Diazodinitn	rophenol		Lead A	zide	Silver Cya	namide	
•	Test No.	1	2	3	4	1	2	ı	2	,
MlAl Squib	Weight - Mgms. Resistance - Ohms Current - Mamps Delay Time - Msecs Flame Duration - Msecs	25. 1.07 250 435.3 26	25 1.00 300 158.07 28	25 1.02 400 23.74 24	25 1.07 500 11.84 24	No Fire No Fire No Fire No Fire No Fire	No Fire No Fire No Fire No Fire No Fire	50 1.02 600	50 1.02 540 15	
	Energy - Ergs	291,000	142,000	38,600	31,500	No Fire	No Fire	Fired	57,654	
	Test No.	1	2			ı	2	1	2	3
T18E4 Carbon Bridge	Volts Condenser - Mfd. Resistance - Ohms Energy - Ergs Remarks	300 .01 1800 0.0 No Fire	300 .1 4200 45,000 Fired			40 .01 1800 0.0 No Fire	50 .01 2400 125 Fired	300 .01 5200 0.0 No Fire	300 1.1 1800 4.95 x 10 ⁵ Fired	300 1.1 25,000 1.28 x 10 ⁹ Fired
	·	I	Cesiu Dinitrob e nz			Pota: Dini: benzof:		Lead Dir	uitrobenzofur	oxan
	Test No.	1	2	3	4	1	2	1	2	3
M1A1 Squib	Weight - Mgms. Resistance - Ohms Current - Mamps Delay Time - Msecs. Flame Duration - Msecs Energy - Ergs.	30 1.12 300 84.51 No Flash 72,000	30 1.07 400 12.34 No Flash 20,000	30 1.08 900 2.14 No Flash 18,700	30 1.05 1000 2.34 No Flash 24,600	16 1.1 300 N.T. Fired N.T.	16 1.1 400 2.8 N.T. 3,000	50 0.91 400 248 82 361,900	50 0.97 550 182 60 582,000	50 1.04 700 196 70 1. x 10 ⁶
	Test No.	1	2	3		1	2	1	2	3
T18 <i>E</i> 4 Carbon Bridge	Volts Condenser - Mfd. Resistance - Ohms Energy - Ergs. Remark	50 .01 1200 0.0 No Fire	75 .01 2100 28 Violent	75 .01 2200 28 Violent		50 .01 2400 0.0 No Fire	100 .01 2200 500 Fired	300 .01 2800 0.0 No Fire	300 1.1 2800 0.0 No Fire	300 .01 2800 450 Fired, No Expl with
				15						10,000 psi
				AANTIN	FA17:41					Par

TABLE 6 (Con't)

	•		Suprous Th	iocyanated Chlorate	•			hiocyanate m Chlorate			ric Thiocya	
	Test No.	1	2	3	4	1	2	3	4	1	2	3
MlAl Squib	Weight - Mgms. Resistance - Ohms Current - Mamps Delay Time - Msecs Flame Duration - Msecs Energy - Ergs	30 1.02 350 58 100+ 72,100	30 1.07 400 23.23 0.5 39,700	30 1.02 500 14.60 0.5 37,200	30 1.09 600 11.55 0.5 45,500	25 1.07 300 31.5 31 53,800	25 1.07 350 36.34 41 47,500	25 1.13 400 24.14 50 43,400	25 1.00 500 18.95 25 47,500	25 1.09 250 56.76 20 39,000	25 1.08 300 24.31 20 23,620	25 1.06 400 16.04 15 27,300
	Test No.	1	2	3		1	2	3	1	2	3	4
T18E4 Carbon Bridge	Volts Condenser - Mfd. Resistance - Ohms Energy - Ergs Remarks	250 .01 1800 0.0 No Fire	300 .01 3200 3500 Fired	300 .01 1900 3500 Fired		300 .01 4800 0.0 No Fire	300 0.11 16,000 49,500 Fired	300 0.11 4800 49,500 Fired	300 .01 42,000 Fired	100 .01 18,000 500 Fired	75 .01 17,000 291.5 No Fire	100 .01 12,000 500 Fired
		Silver DNBF	Cupric DNBF	Mecuric DNBF	Stann MEDN		Cupric MEDNA	Lead MEDNA	Mercury MEDNA			
	Test No.	1	1	1	1	2	1	1	1			
M1A1 Squib	Weight - Mgms. Resistance - Mfd. Current - Mamps. Delay Time - Msecs. Flame Duration - Msecs Energy - Ergs.	No Fire No Fire No Fire No Fire	No Fire No Fire No Fire No Fire No Fire	No Fire No Fire No Fire No Fire No Fire	50 1.1 350 56.03 500+ 35,600	50 1.1 600 37.79 500+ 72,900	No Fire No Fire No Fire No Fire No Fire	No Fire No Fire No Fire No Fire	No Fire No Fire No Fire No Fire No Fire			
	Test No.	1.			1	2	-,					•
T18E4 Carbon Bridge	Volts Condenser - Mfd. Résistance - Ohms Energy - Ergs. Remarks	No Fire No Fire No Fire	No Fire No Fire No Fire	No Fire No Fire No Fire No Fire	300 .01 2400 No Fire	300 1.15 3200 No Fire						



Pir. 1

Picating Arsenal Crime Corps
Apparatus For Determination of Ignition Time and Burning Time and Firing Energy.

A. Ohronograph

A. Chronograph

B. Squib Tester (Firing Apparatus)

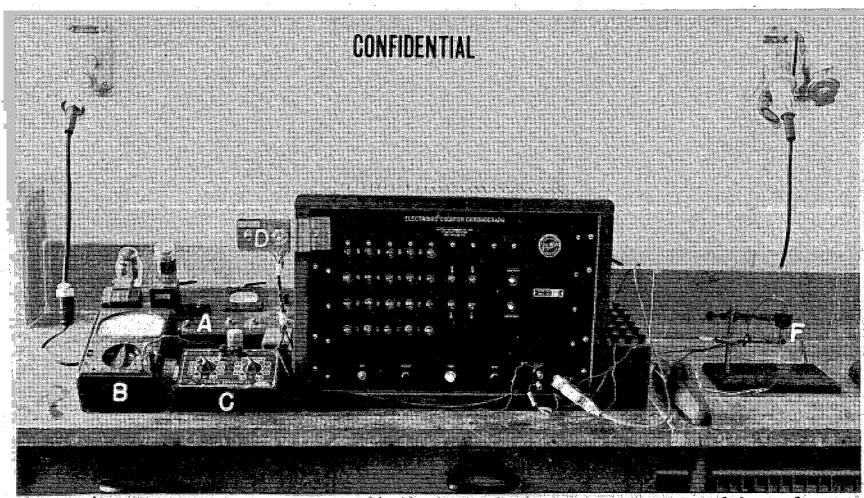
C. Galvanometer

C. Galvanometer

D. Firing leads

C. Galvanometer

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March 1954

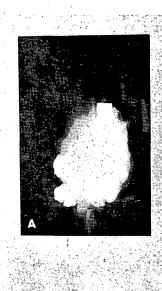
Picetinny Arsenal

Apparatus for Determining Ignition Properties of Explosives on Carbon Bridge Detonators

- A. AC to DC Converter
- OHW Meter
- Variable Condenser

- D. Safety Switch and Firing Switch E. Electronic Counter Chronograph F. Squib Holder Behind Barricade

Figure 2

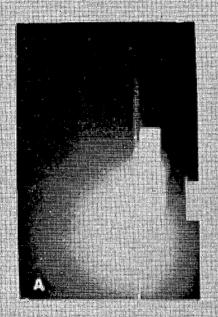


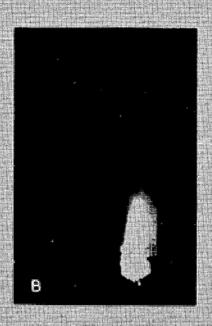






Commence of the contract of th					
M-42137/1		March 1954 PICAT	INNY ARSENAL		ORDNANCE CORPS
704		Flame Charact	eristics of Sta	andard Initiato	rs the
Figure 3		Normal Lead Styphe	ate 💮 💮		
Summer of the grant managers, at 18 19		Colloidal Lead Sty			
	С.	Mercury Fulminate	D. Lead	d Azide	







M-M2131//2

Tables (1934)

FICATINNY ARSENAL

ORDNANCE CORPS

Flame Characteristics of Standard Frimer Mixtures A: PA 100

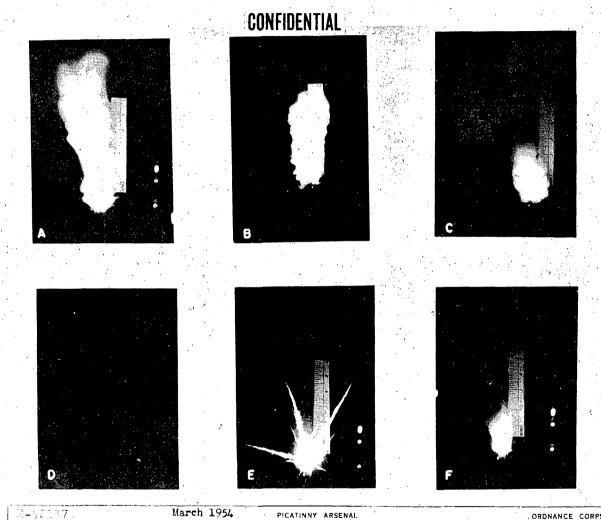
B. Modified 71,

. (66)(je)(r)(jy)(ji)(t

C. FA 70 CONFIDENTIAL

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эn



Flame Characteristics of New Compounds and New Compositions

A. Leas Dinitrobenzfuroxan

B. Cosium Styphnate

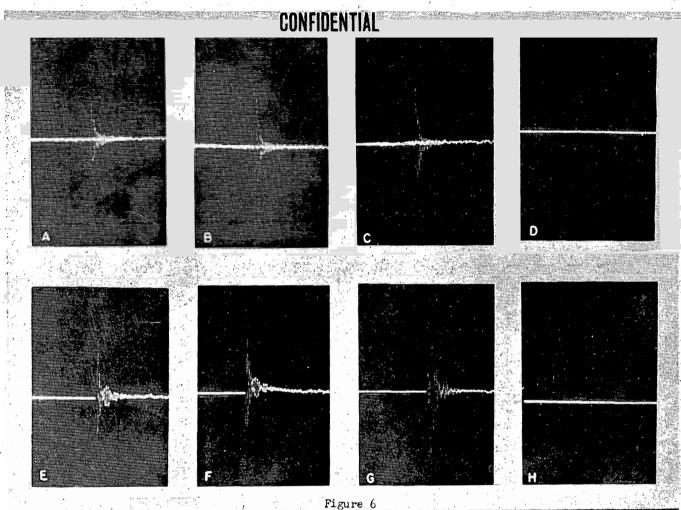
C. Cesium Styphnate

F. Silver Cyanamid

F. Silver Cyanamid

F. Silver Cyanamid

F. Hexamine Chromic Ferchlorate CONFIDENTIAL



M-42139 March 1954 PICATINNY ARSENAL ORDNANCE CORPS Comparison of Shock Waves of Standard Primer Mixtures and Metallic Salts of Dinitro-A. Potassium Dinitrobenzfuroxan E. Basic Lead Styphnate benzfuroxan B. Rubidium "F. NOL 130 Primer (Stab Primer) C. Cesium "G. T61 (Igniter) G. T61 (Igniter) CONFIDENTIAL "CONFIDENTIAL H. FA 70 (Percussion Primer).

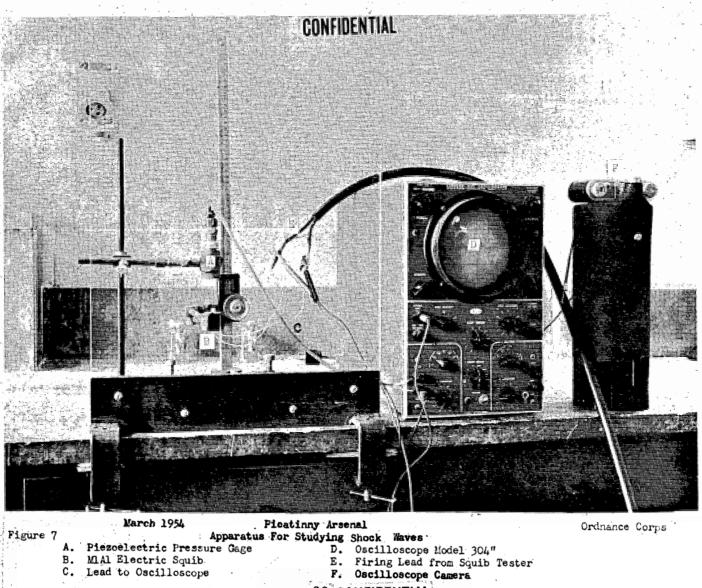


Figure 7

Ordnance Corps



loor Average Particle Size: 192 Microns



THTHACENE 130x Average Particle Size: 190 Microns



SILVER CYANAMIDE Average Particle Size: Agglomorates

Figure 8 CONFIDENTIAL

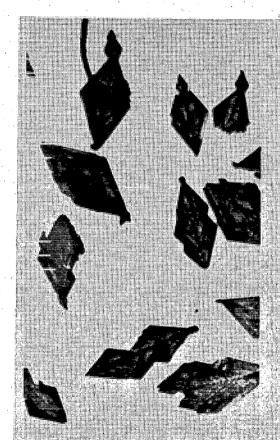
M-42159 March 1955

PICATINNY ARSENAL

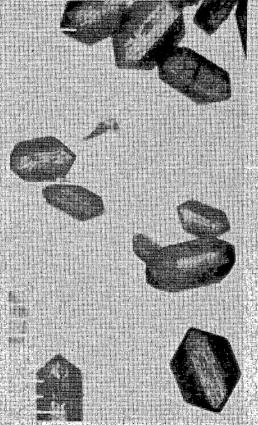
OMDNANCE CORPS

Project No.TA3-5002

Prepared by: A. M. Anzalone Scale: O.1 mm, O.01mm.









NORMAL LEAD STIPHNATZ Average Particle Size:

450m 48 Microne

COLLOIDIAL LEAD STYPHNATE Average Particle Size:

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M-42158

March 1955

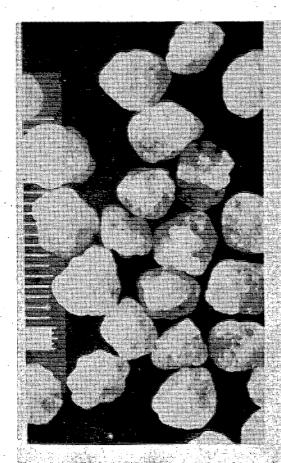
PICATINNY ARSENAL

Pirure 9

ORDHANCE CORPS

Project No. TA3-5002

Prepared by: A. M. Anzalone Scale: 0.1 mm, 0.01 mm



Average Particle Size: 605 Microns



RDX 100x COMPOSITION A_3 20x
Average Particle Size: 388 Microne Average Particle Size: 862 Microne

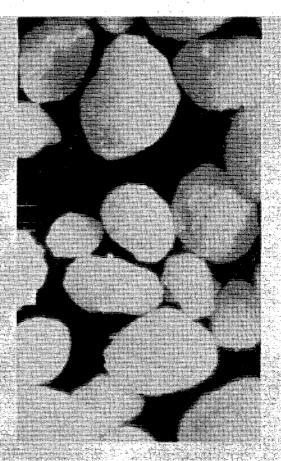


Figure 10 CONFIDENTIAL

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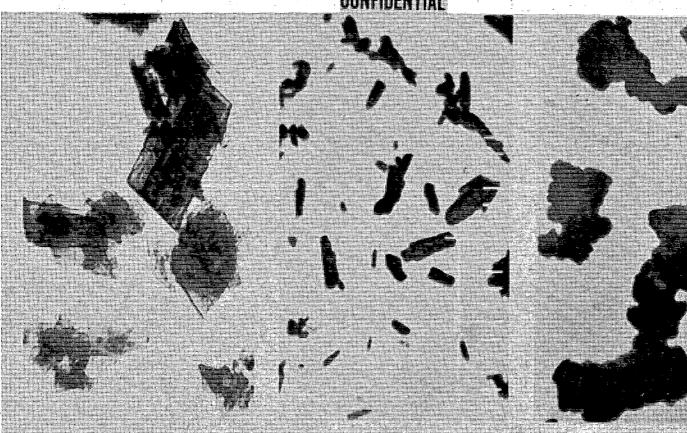
March 1955

PICATINNY ARSENAL

ORDWANCE CORPS

Project No. TA3-5002

Prepared by: A. M. Anzalone Scale: 0.1 mm, 0.01 mm.



POTASSIUM DINITRO BENZO FUROXAN 450x Average Particle Size: 74 Microns

M-42150

D) (AZIO EDIOS (UEZO ESTERIO). Average Particle Size:

500x 28 Migrone

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COMMERCIAL GRADE LEAD AZIDE 520x Average Particle Size: Agglomerates

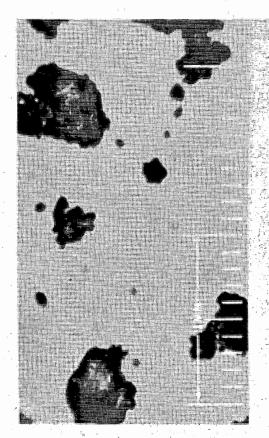
Figure 11

PRETINKS JESENAL

DRONANCE CORPS

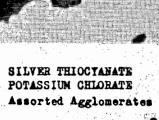
March 1955

Project No. TA3-5002 Prepared by: A. M. Anzalone Scale: 0.1 mm, 0.01 mm.



CUPROUS THIOCYANATE POTASSIUM CHLORATE Asserted Agglomerates

500x



500x

HEXAMINE CHRONIC PERCHLORATE Asserted Agglomerates

500x

Figure 12

M-42161

March 1955

PICATINNY ARSENAL

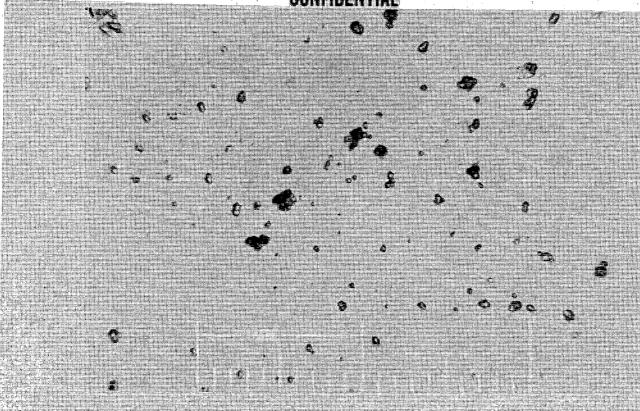
ORDNANCE, CORPS

Project No. TA3-5002

Prepared by: A. M. Anzalone

Scale: 0.1 mm, 0.01mm.

GONHUENHAL



Copper Chlorotetrazole Average Particle Size:

CONFIDENTIAL

500X

∠5 Microns

Figure 13

M-46412 March 1955

PICATINAY ATSENA

ORDNANCE CORPS

Project No. 743-5002

Prepared by: A. M. Anzalone

Scale: 0.1 mm, 0.01 mm.